

## AMENDMENTS TO THE CLAIMS

### **Claims 1 and 2** (previously cancelled)

### **Claim 3** (previously amended)

The method of claim 8, wherein the ketolide is 11,12-dideoxy-3-de[(2,6-dideoxy-3-C-methyl-3-0-methyl-alpha-6-ribohexopyranosyl)oxy]-6-0-methyl-3-oxo-12,11-([oxycarbonyl [[[2-[4-(3-pyridinyl)-1H-imidazol-1-yl]ethoxy]-methyl]imino]]erythromycin.

### **Claim 4** (previously amended)

The method of claim 8, wherein the ketolide is 11,12-dideoxy-3-de((2,6-dideoxy-3-C-methyl-3-0-methyl-alpha-L-ribohexopyranosyl)oxy)-6-0-methyl-3-oxo-12,11-(oxycarbonyl((4-(3-(3-pyridinyl)-1H-1,2,4-triazol-1-yl)butyl)imino)erythromycin.

### **Claim 5** (previously amended)

The method of claim 8, wherein the ketolide is 11,12-dideoxy-3-de((2,6-dideoxy-3-C-methyl-3-0-methyl-alpha-L-ribohexopyranosyl)oxy)-2-fluoro-6-0-methyl-3-oxo-12,11-(oxycarbonyl((4-(4-(3-pyridinyl)-1H-imidazol-1-yl)butyl)imino))erythromycin (A isomer).

**Claim 6** (previously amended)

The method of claim 8, wherein the ketolide is 11,12-dideoxy-3-de((2,6-dideoxy-3-C-methyl-3-O-methyl- $\alpha$ -L-ribohexopyranosyl)oxy)-6-O-methyl-3-oxo-12,11-(oxycarbonyl((4-(4-(3-pyridinyl)-1H-imidazol-1-yl)butyl)imino))erythromycin.

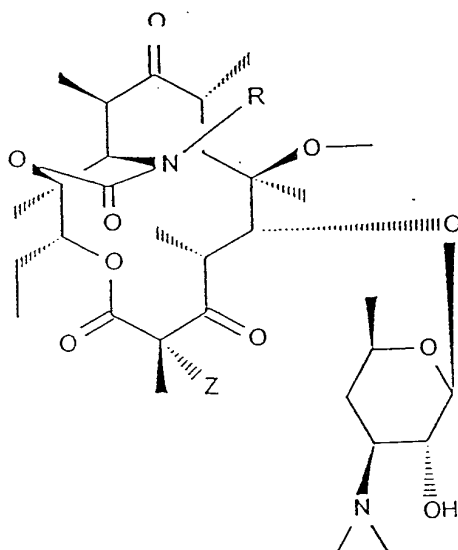
**Claim 7** (previously cancelled)

**Claim 8** (currently amended)

A method of treating atherosclerosis ~~preventing the arterial complications~~ in a patient comprising selecting a patient with atherosclerosis ~~arterial complications~~ and administering to said patient an effective amount of a ketolide or its non-toxic, pharmaceutically acceptable acid addition salts sufficient to treat atherosclerosis ~~prevent arterial complications~~ in said patient.

**Claim 9** (previously added)

The method of claim 8 wherein the ketolide has the formula



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wherein R is  $-(CH_2)_mO_n(X)YAr$ , m and n are individually 0 or 1, X is selected from the group consisting of  $-(NH)_a-$ ,  $-CH_2-$  and  $-SO_2-$ , a is 0 or 1, Y is  $-(CH_2)_b-(CH=CH)_c-(CH_2)_d-$ , c is 0 or 1,  $b + c + d \leq 8$ , Z is hydrogen or halogen and Ar is unsubstituted or substituted aryl or heteroaryl.

**Claim 10** (previously added)

The method of claim 8 wherein the ketolide is orally administered at 50 to 500 mg per day.